

We claim:

1. An immunogenic fusion protein comprising (a) a modified NS3 polypeptide comprising at least one amino acid substitution to the HCV NS3 region, such that  
5 protease activity is inhibited, and (b) at least one polypeptide derived from a region of the HCV polyprotein other than the NS3 region.
2. The fusion protein of claim 1, wherein the modification comprises a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165,  
10 numbered relative to the full-length HCV-1 polyprotein.
3. The fusion protein of claim 1, wherein the protein comprises a modified NS3 polypeptide, an NS4 polypeptide, an NS5a polypeptide, and optionally a core polypeptide.  
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4. The fusion protein of claim 3, wherein the protein further comprises an NS5b polypeptide, and optionally a core polypeptide.
5. The fusion protein of claim 3, wherein the protein further comprises an E2  
20 polypeptide, a p7 polypeptide, an NS2 polypeptide, and optionally a core polypeptide.
6. The fusion protein of claim 3, wherein the protein further comprises an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, and optionally a core polypeptide.  
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7. The fusion protein of claim 3, wherein the protein further comprises an E2 polypeptide, and optionally a core polypeptide.
8. The fusion protein of claim 3, wherein the protein further comprises an E1  
30 polypeptide, an E2 polypeptide, and optionally a core polypeptide.

9. The fusion protein of claim 1, wherein the protein comprises an E2 polypeptide, a modified NS3 polypeptide, and optionally a core polypeptide.

5           10. The fusion protein of claim 1, wherein the protein comprises an E1 polypeptide, an E2 polypeptide, a modified NS3 polypeptide, and optionally a core polypeptide.

10           11. The fusion protein of claim 1, wherein the polypeptides of (a) and (b) are derived from the same HCV isolate.

12. The fusion protein of claim 1, wherein at least one of the polypeptides present in the fusion is derived from a different isolate than the modified NS3 polypeptide.

15           13. An immunogenic fusion protein consisting essentially of, in amino terminal to carboxy terminal direction:

          (a) a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, and an NS5a polypeptide;

          (b) a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide and an NS5b polypeptide;

25           (c) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, and an NS5a polypeptide;

30           (d) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to

His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, and an NS5a polypeptide;

5 (e) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide and an NS5b polypeptide;

10 (f) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide and an NS5b polypeptide;

15 (g) an E2 polypeptide and a modified NS3 polypeptide comprising substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited;

20 (h) an E1 polypeptide, an E2 polypeptide and a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited;

(i) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide and a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited; or

25 (j) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide and a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited.

14. An immunogenic fusion protein consisting essentially of, in amino terminal to carboxy terminal direction:

(a) a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide, and a core polypeptide;

(b) a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide, an NS5b polypeptide and a core polypeptide;

(c) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide and a core polypeptide;

(d) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide and a core polypeptide;

(e) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide, an NS5b polypeptide and a core polypeptide;

(f) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, an NS4 polypeptide, an NS5a polypeptide, an NS5b polypeptide and a core polypeptide;

(g) an E2 polypeptide, a modified NS3 polypeptide comprising substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, and a core polypeptide;

5 (h) an E1 polypeptide, an E2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, and a core polypeptide;

(i) an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3  
10 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, and a core polypeptide; or

(j) an E1 polypeptide, an E2 polypeptide, a p7 polypeptide, an NS2 polypeptide, a modified NS3 polypeptide comprising a substitution of an amino acid corresponding to  
15 His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-length HCV-1 polyprotein such that protease activity is inhibited, and a core polypeptide.

15. A modified NS3 polypeptide comprising a substitution of an amino acid corresponding to His-1083, Asp-1105 and/or Ser-1165, numbered relative to the full-  
20 length HCV-1 polyprotein such that protease activity is inhibited when the modified NS3 polypeptide is present in an HCV fusion protein.

16. A composition comprising an immunogenic fusion protein according to claim 1 in combination with a pharmaceutically acceptable excipient.

25 17. A composition comprising an immunogenic fusion protein according to claim 13 in combination with a pharmaceutically acceptable excipient.

18. A composition comprising an immunogenic fusion protein according to claim 14 in combination with a pharmaceutically acceptable excipient.

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19. A method of stimulating a cellular immune response in a vertebrate subject comprising administering a therapeutically effective amount of the composition of claim 16.

5           20. A method of stimulating a cellular immune response in a vertebrate subject comprising administering a therapeutically effective amount of the composition of claim 17.

21. A method of stimulating a cellular immune response in a vertebrate subject  
10 comprising administering a therapeutically effective amount of the composition of claim 18.

22. A method for producing a composition comprising combining the  
immunogenic fusion protein of claim 1 with a pharmaceutically acceptable excipient.  
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23. A method for producing a composition comprising combining the  
immunogenic fusion protein of claim 13 with a pharmaceutically acceptable excipient.

24. A method for producing a composition comprising combining the  
20 immunogenic fusion protein of claim 14 with a pharmaceutically acceptable excipient.

25. A polynucleotide comprising a coding sequence encoding a fusion protein  
according to claim 1.

25           26. A polynucleotide comprising a coding sequence encoding a fusion protein  
according to claim 13.

27. A polynucleotide comprising a coding sequence encoding a fusion protein  
according to claim 14.  
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28. A polynucleotide comprising a coding sequence encoding a polypeptide according to claim 15.

5           29. A recombinant vector comprising:  
          (a) the polynucleotide of claim 25; and  
          (b) at least one control element operably linked to said polynucleotide, whereby said coding sequence can be transcribed and translated in a host cell.

10           30. A recombinant vector comprising:  
          (a) the polynucleotide of claim 26; and  
          (b) at least one control element operably linked to said polynucleotide, whereby said coding sequence can be transcribed and translated in a host cell.

15           31. A recombinant vector comprising:  
          (a) the polynucleotide of claim 27; and  
          (b) at least one control element operably linked to said polynucleotide, whereby said coding sequence can be transcribed and translated in a host cell.

20           32. A recombinant vector comprising:  
          (a) the polynucleotide of claim 28; and  
          (b) at least one control element operably linked to said polynucleotide, whereby said coding sequence can be transcribed and translated in a host cell.

25           33. A host cell comprising the recombinant vector of claim 29.

          34. A host cell comprising the recombinant vector of claim 30.

          35. A host cell comprising the recombinant vector of claim 31.

30           36. A host cell comprising the recombinant vector of claim 32.

37. A method for producing an immunogenic fusion protein, said method comprising culturing a population of host cells according to claim 33 under conditions for producing said protein.

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38. A method for producing an immunogenic fusion protein, said method comprising culturing a population of host cells according to claim 34 under conditions for producing said protein.

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39. A method for producing an immunogenic fusion protein, said method comprising culturing a population of host cells according to claim 35 under conditions for producing said protein.

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40. A method for producing a polypeptide, said method comprising culturing a population of host cells according to claim 36 under conditions for producing said polypeptide.